

1
2 **Claims**
3

4 What we claim is:

5
6 sub a' 1. An endosomal lysing agent comprising a compound having one or more hydrolyzable
7 functional moieties and wherein said compound is capable of effecting the lysis of an endosome
8 in response to a change in pH.
9

10 2. The endosomal lysing agent of claim 1, comprising a biocompatible compound.
11

12 sub b1 3. The endosomal lysing agent of claim 1, comprising a biodegradable compound.
13

14 4. The endosomal lysing agent of claim 1, comprising a biocompatible and biodegradable
15 compound.
16

17 sub a' 5. An endosomal lysing agent comprising a compound having one or more hydrolyzable
18 functional moieties and one or more ionizable functional moieties, and wherein said compound
19 is capable of effecting the lysis of an endosome in response to a change in pH.
20

21 6. The endosomal lysing agent of claim 5, comprising a biocompatible compound.
22

23 sub b1 7. The endosomal lysing agent of claim 5, comprising a biodegradable compound.
24

25 8. The endosomal lysing agent of claim 5, comprising a biocompatible and biodegradable
26 compound.
27

28 9. The endosomal lysing agent of claim 1, 2, 3, 4, 5, 6, 7, or 8 comprised of a polymer.
29

1 10. The endosomal lysing agent of claim 9, wherein the hydrolysis of said one or more
2 hydrolyzable functional moieties effects a hydrophobic/hydrophilic transition of said compound.

3
4 11. The endosomal lysing agent of claim 10, wherein said hydrolysis further effects the
5 release of a compound capable of disrupting lipid bilayers.

6
7 12. The endosomal lysing agent of claim 5, wherein said one or more ionizable functional
8 moieties comprises proton acceptor sites.

9
10 13. The endosomal lysing agent of claim 1 or 5, wherein said one or more hydrolyzable
11 functionalities is independently selected from the group consisting of ortho-ester, hydrazone, and
12 cis-acetonyl

13
14 14. The endosomal lysing agent of claim 13, wherein each of said ortho-ester containing
15 monomers is selected from the group consisting of N-[2-methyl-1,3-O-ethoxyethylidene-
16 prpanediol]methacrylamide, ortho-ester derivatives of tartaric acid, ortho-ester derivatives of
17 treitol, and ortho-ester derivatives of dithiothreitol.

18
19 15. The polymeric lysing agent of claim 9, wherein the polymeric lysing agent is combined
20 in a form selected from the group consisting of:

21 mixed polymers;

22 linear co-polymers;

23 branched co-polymers; and

24 dendrimer branched co-polymers.

25
26 16. The lysing agent of claim 9, wherein said agent is further functionalized with a targeting
27 agent selected from the group consisting of low density lipoproteins, transferrin,
28 asiaglycoproteins, gp120 envelope protein of human immunodeficiency virus, antibodies and
29 carbohydrates.

Sub 24

17. A biocompatible composition comprising:
a packaging agent, characterized by an ability to bind to a therapeutic agent and mediate import into endosomes; and
a lysing agent comprising a compound having one or more hydrolyzable functional moieties and wherein said compound is capable of effecting the lysis of an endosome in response to a change in pH.

Sub B3) 18. The biocompatible composition of claim 17, wherein said compound further comprises one or more ionizable functional moieties.

19. The biocompatible composition of claim 17 or claim 18, wherein said composition comprises a polymer.

20. The biocompatible composition of claim 17 or 18, wherein said packaging agent and said lysing agent are combined in a form selected from the group consisting of:

Sub B4)
mixed polymers;
linear co-polymers;
branched co-polymers; and
dendrimer branched co-polymers.

21. The biocompatible composition of claim 17 or claim 18, wherein said therapeutic agent comprises a nucleic acid.

22. The biocompatible composition of claim 17 or claim 18, wherein the packaging agent associates with the therapeutic agent through a covalent interaction.

23. The biocompatible composition of claim 17 or claim 18, wherein the packaging agent associates with the therapeutic agent through a non-covalent interaction.

1 24. The composition of claim 17 or claim 18, wherein the packaging agent condenses the
2 nucleic acid.

3
4 25. The composition of claim 17 or claim 18, wherein the packaging agent condenses the
5 nucleic acid to a size less than 150 nm.

6
7 26. The composition of claim 17 or claim 18, wherein the packaging agent comprises a
8 material with high charge density.

9
10 27. The composition of claim 26, wherein said packaging agent comprises a tertiary amine or
11 a quaternary amine.

12
13 28. The composition of claim 27, wherein said packaging agent is selected from the group
14 consisting of 2-[dimethylamino]ethyl methacrylate, (3-aminopropyl)methacrylamide, 2-
15 aminoethyl methacrylamide, aspartic acid, glutamic acid and polymers thereof.

16
17 29. The composition of claim 17 or claim 18, wherein the hydrolysis of said one or more
18 hydrolyzable functional moieties effects a hydrophobic/hydrophilic transition of said compound.

19
20 30. The composition of claim 17 or claim 18, wherein said hydrolysis further effects the
21 release of a compound capable of disrupting lipid bilayers.

22
23 31. The composition of claim 18, wherein said one or more ionizable functional moieties
24 comprises proton acceptor sites.

25
26 ^{sub 25} 32. A cell delivery composition comprising:
27 a compound to be delivered to a cell;
28 a delivery agent bound to the compound; and

1 an endosomolytic agent comprising a compound capable of effecting the lysis of an
2 endosome in response to a change in pH.

3
4 33. The cell delivery composition of claim 32, wherein said endosomolytic agent comprises a
5 compound having one or more hydrolyzable functionalities.

6
7 34. The cell delivery composition of claim 32, wherein said endosomolytic agent comprises a
8 compound having one or more hydrolyzable functionalities and one or more ionizable
9 functionalities.

10
11 35. The cell delivery composition of claim 32, wherein the compound to be delivered to a
12 cell is selected from the group consisting of anti-AIDS substances, anti-cancer substances,
13 antibiotics, immunosuppressants, anti-viral substances, enzyme inhibitors, neurotoxins, opioids,
14 hypnotics, antihistamines, lubricants, tranquilizers, anti-convulsants, muscle relaxants, anti-
15 Parkinson substances, anti-spasmodics and muscle contractants, miotics, anti-cholinergics, anti-
16 glaucoma compounds, anti-parasite compounds, anti-protozoal compounds, anti-hypertensives,
17 analgesics, anti-pyretics, anti-inflammatory agents, local anesthetics, ophthalmics,
18 prostaglandins, anti-depressants, anti-psychotic substances, anti-emetics, imaging agents,
19 specific targeting agents, neurotransmitters, proteins, cell response modifiers, vaccines, anti-
20 sense agents, RNA and ribozymes.

21
22 36. A non-immunogenic artificial virus less than 150 nm in size, comprising:
23 a nucleic acid packaging agent;
24 an endosomal lysing component capable of effecting the lysis of an endosome in
25 response to a change in pH; and
26 a nucleic acid.

27
28 37. The artificial virus of claim 36, wherein said endosomal lysing component comprises a
29 compound having one or more hydrolyzable functionalities.

1 38. The artificial virus of claim 36, wherein said endosomal lysing component comprises a
2 compound having one or more hydrolyzable functionalities and one or more ionizable
3 functionalities.

4
5 ^{Sub 26} 39. A method of lysing an endosome, the method comprising the steps of:
6 providing a composition for endosomal uptake into the cell; and
7 contacting the composition with the cell in the presence of an endosomal lysing agent
8 capable of effecting the lysis of an endosome in response to a change in pH.

9
10 40. The method of claim 39, wherein said endosomal lysing agent comprises a compound
11 having one or more hydrolyzable functionalities.

12
13 ^{Sub 28} 41. The method of claim 39, wherein said endosomal lysing agent comprises a compound
14 having one or more hydrolyzable functionalities and one or more ionizable functionalities.

15
16 ^{Sub 29} 42. A method for introducing a therapeutic agent into a cell or a subcellular component, the
17 method comprising the steps of:
18 providing a biocompatible delivery composition comprising:
19 a packaging agent;
20 an endosomal lysing component capable of effecting the lysis of an endosome in
21 response to a change in pH; and
22 a nucleic acid; and
23 contacting the composition with cells.

24
25 43. The method of claim 42, wherein said endosomolytic agent comprises a compound
26 having one or more hydrolyzable functionalities.

27
28 44. The method of claim 42, wherein said endosomolytic agent comprises a compound
29 ^{Sub 29} having one or more hydrolyzable functionalities and one or more ionizable functionalities.

1 45. The method of claim 42, further comprising contacting the composition with cells in the
2 absence of a known endosomal lysing component selected from the group consisting of
3 chloroquine, polyethyleneimine, fusogenic peptides, inactivated adenoviruses and combinations
4 thereof.

APP 310)